Scapania paraphyllia, a new synonym of Scapania koponenii (Marchantiophyta, Scapaniaceae) with special reference to paraphyllia, pseudoparaphyllia, and paraphyses

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Scapania paraphyllia T.Cao, C.Gao, J.Sun & B.R.Zuo and *S. koponenii* Potemkin are both endemic to China. They share considerable morphological similarities except that the former species has pseudoparaphyllia. In order to elucidate the taxonomic status of the two species, we investigated morphological characters possessed by their type specimens and additional collections, and reconstructed a species phylogeny for *S. koponenii*, *S. paraphyllia*, and closely related species. The phylogenetic trees were constructed using maximum parsimony, maximum likelihood and Bayesian inference analyses based on the nuclear marker nrITS and the plastid markers *trn*L-F region and *atp*B-*rbc*L spacer. Both morphological and molecular evidence contradicted the hypothesis that *S. paraphyllia* and *S. koponenii* are separate species. The pseudoparaphyllia reported for *S. paraphyllia* are confirmed as paraphyses. The paraphyllia, paraphyses, and pseudoparaphyllia in *Scapania* are discussed.

Keywords: China, Hepaticae, Liverwort, Molecular phylogeny, Taxonomy

Introduction

Scapania (Dumort.) Dumort., with 100 currently accepted species (Söderström et al., 2016), is the largest genus in its own family Scapaniaceae, most species of which occur in northern temperate regions. The genus is well characterised and easily recognised by the (1) lack of underleaves, (2) prominently lateral-intercalary branching, (3) leaves complicatebilobed, with ventral leaf lobes (also called leaf lobules) often rotund to ovate or even lingulate and larger than dorsal leaf lobes (also called leaf lobes), (4) presence of a usually winged keel, (5) leafmargins usually denticulate, occasionally entire, (6) cuticles smooth to coarsely verruculose or obviously verrucose, and (7) dorso-ventrally compressed perianths, usually with truncate and entire mouths. Scapania was extensively studied using both morphological (Müller, 1905; Amakawa & Hattori, 1953, 1955; Amakawa, 1964; Schuster, 1974; 1954. Potemkin, 1998, 1999a, 1999b, 2002) and molecular data to resolve species delimitations and supraspecific classifications (Vilnet *et al.*, 2010; Heinrichs *et al.*, 2012).

So far, about 50 Scapania species have been recognised in China (Cao & Sun, 2008; Jia & He, 2013). Among them, Scapania paraphyllia T.Cao, C.Gao, J.Sun & B.R.Zuo is a species endemic to China, and known only from the two type specimens found in Zhejiang province (Zuo et al., 2007; Jia & He, 2013). This species is characterised by the keel being about 1/4-1/3 the length of the leaf lobule, cuticle of leaves verrucose with large papillae, and shoots with pseudoparaphyllia in the leaf axils or on stems (Zuo et al., 2007). During recent studies of Scapania in China, we observed that S. paraphyllia shared numerous common characters with S. koponenii Potemkin, another endemic species of China, previously known from south-eastern China (Potemkin, 2000). It seemed that the presence of 'pseudoparaphyllia' was the only character differing between these two species.

This study aimed to test the circumscription of *S. paraphyllia* and *S. koponenii* by conducting phylogenetic analyses based on 38 samples including

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S. paraphyllia and *S. koponenii* from their type localities and neighbouring provinces of China using nuclear and plastid DNA regions (nrITS, *trn*L-F region, and *atp*B-*rbc*L spacer). In conclusion, we propose a new synonym in accordance with the evidence from the morphological characters and molecular topologies.

Materials and Methods

Taxon sampling

To assess the phylogenetic relationship between S. paraphyllia and S. koponenii, we used a dataset consisting of 38 samples from 15 species in eight sections belonging to subg. Scapania. Among them, the sequences of 11 samples were newly obtained and those of the remaining 27 samples were downloaded from GenBank (http://www.ncbi.nlm.nih.gov/ genbank/). According to the phylogenies generated by Heinrichs et al. (2012) and Váňa et al. (2012), S. aequiloba (Schwagr.) Dumort. and S. curta (Mart.) Dumort., representing the sections Aequilobae and *Curtae*, respectively, were chosen as the outgroup in this study. Voucher specimens for the 11 new samples were carefully examined by light microscopy (Olympus BX 43). Voucher information and GenBank accession numbers for all sequences used in the current study are listed in Appendix 1.

DNA extraction, PCR amplification, and sequencing

Tissue from the distal parts of gametophyte shoots was isolated from herbarium specimens collected within the last three years. Total genomic DNA was extracted with Qiagen DNeasy Plant® Mini Kit. Primers for PCR were those described in previous studies: trnL-F region from Taberlet et al. (1991), atpB-rbcL spacer from Feldberg et al. (2010), and nrITS region from Groth et al. (2003). PCR amplification was carried out as follows. For trnL-F and atpB-rbcL, an initial denaturation at 94°C for 4 min was followed by 33 cycles of 1 min denaturation at 94°C, 1 min annealing at 51°C (trnL-F) or 50°C (atpB-rbcL), and 1 min elongation at 72°C, then a final elongation step of 10 min at 72°C. For nrITS, an initial denaturation step at 94°C for 4 min was followed by 33 cycles of 1 min denaturation at 94°C, 1 min annealing at 51°C and 1.5 min elongation at 72°C, then a final 10 min elongation at 72°C. The PCR products were first tested by electrophoresis in an agarose gel to confirm successful amplification. The successful amplifications were sent to Shanghai Majorbio Bio-Pharm Technology Co. Ltd., China (http://www.majorbio .bioon.com.cn) for bidirectional sequencing with the same primers as used in PCR.

Phylogenetic analysis

In the phylogeny analyses, 108 sequences were included in the final dataset, of which 32 sequences were newly generated, whilst 76 were obtained from Genbank. All sequences were assembled and manually aligned in PhyDE®0.9971 (Müller *et al.*, 2005). Ambiguous positions in the dataset were excluded from subsequent phylogenetic analyses; the gaps and missing nucleotides were scored as missing data. Maximum parsimony (MP), maximum likelihood (ML), and Bayesian inference (BI) phylogenetic methods were employed to infer phylogenies.

Maximum parsimony analyses were conducted in PAUP 4.0b10 (Swofford, 2003). Command files based on the parsimony ratchet (Nixon, 1999) were generated by the programme PRAP2 (Müller, 2007). All characters were equally weighted and treated as unordered and heuristic searches were performed for 10,000 replicates with tree bisection and reconnection (TBR) branch swapping. The three single locus datasets were first subjected to separate MP analyses to examine for possible incongruence. The 70% bootstrap criterion was used to identify incongruent signals (Mason-Gamer & Kellogg, 1996) by visual comparisons of the tree topologies and support values. No evidence of incongruence was shown and thus the three datasets were combined for subsequent phylogenetic analyses. Maximum likelihood analyses were conducted using RAxML8.2 (Stamatakis, 2014). A rapid bootstrap analysis was run using the GTRGAMMAI substitution model with 1000 replicates. Partitioned BI analyses were performed in MrBayes v.3.0 (Huelsenbeck & Ronquist, 2001). The optimal nucleotide substitution models for each region: HKY + G (trnL-F), GTR + I (atpB-rbcL), and GTR + I + G (nrITS) were calculated using MrModeltest v.2.3 (Nylander, 2004) as implemented in PAUP* v.4.0b10 and based on the Akaike Information Criterion (AIC). The data were analysed using Markov Chain Monte Carlo (MCMC), performing four parallel analyses with four chains each run for 2 million generations, and sampling trees every 1000 generations. The first 10% of the sampled trees were discarded as burn-in determined by Tracer v1.5 (Rambaut & Drummond, 2009). From the remaining tree set, a single majority-rule consensus tree with the posterior probabilities (PP) was generated. Trees were depicted in TreeGraph v.2 (Stöver & Müller, 2010).

Results

The combined alignment consists of 1698 characters of 38 samples, including 377 in the locus trnL-F, 595 in atpB-rbcL and 726 in nrITS. Of the total characters, 1375 were constant, 228 were parsimony informative, 95 were variable but parsimony uninformative. The



Figure 1 Majority-rule consensus tree based on Bayesian analyses of the combined dataset of *trn*L-F, *atp*B-*rbc*L, and nrITS. Bayesian posterior probability values (PP) followed by bootstrap percentage values from maximum likelihood (MLBS) and maximum parsimony (MPBS) are indicated at branches (from left to right: PP/MLBS/MPBS). Branches in bold are highly supported by all analytical methods (PP \ge 0.99; MLBS \ge 95; MPBS \ge 85). Support values of PP < 0.95, MLBS < 70, and MPBS < 70 are not shown or indicated by a dash (-).

MP analyses generated four maximally parsimonious trees by a length of 525 steps, with a consistency index (CI) of 0.749 and a retention index (RI) of 0.884. BI and ML analyses resulted in topologies congruent with the topology from MP analyses. Thus the consensus tree from the Bayesian inference analyses is presented in Figure 1, with posterior probabilities (PP \ge 0.95) indicated on the branches as well as bootstrap support values (BS \ge 70) obtained from the likelihood and parsimony analyses separately.

In the phylogenetic topology (Figure 1), the ingroup contained 13 species. Nine species with multiple accessions were resolved in monophyletic lineages (PP = 1; MLBS \geq 97; MPBS \geq 87), while only *S. ciliata* Sande Lac. was moderately supported in both the ML and MP analyses (MLBS = 82; MPBS = 74). The 13 species represent six sections of the Heinrichs *et al.* (2012) phylogeny of *Scapania* (Figure 1). In sect. *Ciliatae*, nine samples of *S. koponenii* with two samples of *S. paraphyllia* were grouped together to form a clearly

monophyletic clade with strong support (PP = 1; MLBS = 100; MPBS = 100).

Discussion

Scapania paraphyllia was described based on two herbarium specimens (Z.L. Liu 554, holotype; Z.L. Liu 553, paratype) originally determined as S. stephanii Müll.Frib. from Zhejiang province (Zuo et al., 2007). Our examination reveals that both holotype and paratype contain both male and female plants, and that the so-called 'pseudoparaphyllia' in S. paraphyllia are actually in male bract-axils or occasionally on stems just out of the bract-base of androecia (Figure 2D, N). The paraphyses in S. paraphyllia' because pseudoparaphyllia are restricted to the areas of the stem around branch primordia, and not associated with antheridia and archegonia (Magill, 1990).

Scapania koponenii was originally described by Potemkin (2000) based on samples from five provinces of China, including Fujian, Guangdong, Hunan,



Figure 2 Scapania koponenii Potemkin. (A) Portion of shoot, ventral view. (B) Portion of shoot, dorsal view. (C) Portion of shoot with perianth, ventral view. (D) Portion of shoot with androecia, ventral view. (E) Leaf. (F, O, P) Male bracts. (G) Apical portion of leaf lobule. (H) Gemmae. (I) Sector of perianth mouth. (J) Cross section of stem. (K, Q, R) Paraphyses. (L) Median cells of leaf lobule. (M) Papillae of leaf lobule. (N) Portion of male plant, showing antheridia and paraphyses in male bracts. A–J, L–N, and Q from *Zhang et al.* 20150830-178A (HSNU), K and O from *Li* 95233 (paratype of *S. koponenii*, HSNU), P and R from *Liu* 554 (holotype of *S. paraphyllia*, HSNU).

Jiangxi, and Zhejiang. It has thus far only been found in China, but does not seem rare. It is widely distributed in Fujian, Guangdong, Guangxi, Guizhou, Hunan, Jiangxi, and Zhejiang (Zhu *et al.*, 2006; Cao & Sun, 2008); it mainly grows on wet cliffs, wet rocks or rocks with a thin layer of soil, usually in clumps in moist habitats at altitudes of 400–1800 m. This species is well characterised and easily recognised



Figure 3 *Scapania koponenii* Potemkin. Portion of male plant, showing antheridia associated with paraphyses. Drawn from *Zhang et al.* 20150830-178A (HSNU).

by the small plants (mostly 1–2 cm long) (Figure 2A, B), the leaf lobe about 1/3-1/2 the size of leaf lobule, the keel about 1/4-2/5 the length of the leaf lobule (Figure 2E), the irregular marginal toothed leaf and perianth mouth (Figure 2C, G, and I), and the cuticle densely to coarsely papillose (Figure 2M). In addition, the androecia of this species are short with subequally bilobed male bracts in 2-3 pairs with a longer keel (Potemkin, 2000) (Figure 2D, F, and O). Male plants of numerous specimens contain numerous paraphyses which are the same as those in the type specimens of S. paraphyllia (Figure 2D, K, N, Q, and R; Figure 3). Morphological observations on S. koponenii and S. paraphyllia reveal that the two taxa have no significant differences as suggested by our molecular analyses. Scapania paraphyllia is thus proposed as a new synonym of S. koponenii.

In bryophytes paraphyses are hyaline or yellowish, usually uniseriate, hairs often associated with antheridia and archegonia (Magill, 1990; Merced-Alejandro & Sastre-De Jesús, 2009). Back in 1905, paraphyses were reported in Scapania americana Mull.Frib., S. aspera M.Bernet & Bernet, S. compacta (Roth) Dumort., S. ligulata subsp. stephanii (Mull.Frib.) Potemkin, Piippo & T.J.Kop., S. irrigua (Nees) Nees, S. kaurinii Ryan., S. secunda Steph., S. umbrosa (Schrad.) Dumort., and S. undulata (L.) Dumort. (Müller, 1905). Schuster (1974) and Amakawa (1964, 1967, 1968) also described paraphyses in further species, e.g., S. brevicaulis Taylor, S. curta, S. hyperborea Jørg., S. komagadakensis Amakawa, S. nemorea (L.) Grolle (as S. nemorosa (L.) Dumort.), S. nipponica (Amakawa & S.Hatt.) Amakawa, S. obcordata (Berggr.) S.W.Arnell., S. okamurana Steph. ex Amakawa & S.Hatt. (= S. parvitexta Steph.). Recently Potemkin (2000, 2001) also reported the occurrence of paraphyses in S. davidii Potemkin, S. koponenii Potemkin and S. sinikkae Potemkin. The paraphyses reported in all species mentioned above are associated with antheridia, except for S. davidii whose paraphyses also occur in female bracts (Majumdar et al., 2016). Paraphyses are variable in size and shape in *Scapania*. They are usually linear, lanceolate, subulate, lamellate, and foliaceous and their margins are entire, or densely ciliate to dentate (Schuster, 1974; Majumdar et al., 2016). In Scapania koponenii, the paraphyses are yellowish to reddish-brown, linear to lanceolate, 21-128×188- $562 \,\mu\text{m}$, entire or occasionally with teeth at the margin (Figure 2K, N, Q, R; Figure 3). Although reduced paraphyses were treated as advanced morphological characters in Scapania (Potemkin, 2002), the taxonomic value of paraphyses still needs to wait for further studies.

The word pseudoparaphyllium was first used by Warnstorf (1904-1906) who defined it as a foliose structure around the branch primordia in Rhynchostegium confertum (Dicks.) Schimp. (a moss). The term, however, was later confused (Ireland, 1971; Schofield & Hebant, 1984), and it was difficult to separate from the term paraphyllium. Ignatov & Hedenäs (2007) explored the homologies of pseudoparaphyllia in pleurocarpous mosses and suggested that the term pseudoparaphyllium be used for any structures developed near or at the base of branch primordia. In Scapania pseudoparaphyllia have been reported in only two species, S. macroparaphyllia T.Cao C.Gao & J.Sun (Cao et al., 2004) and S. paraphyllia (Zuo et al., 2007). In the latter the pseudoparaphyllia proved to be mis-interpreted paraphyses. Owing to the limited samples available in the present study it remains unknown whether same mis-understanding has occured in the S. macroparaphyllia.

Paraphyllia are tiny filaments, scales, or leaf-like structures scattered on the stems of some leafy liverworts and pleurocarpous mosses (Malcolm & Malcolm, 2006). Although paraphyllia are not frequent in liverworts, they have been reported and used as one of the key characters in the taxonomy of Plagiochila (Dumort.) Dumort. (Grolle & So, 1999; So, 2001). In Scapania, paraphyllia have been known in only a few species, e.g., S. bolanderi Austin, S. maxima Horik., S. ornithopodioides (With.) Waddell, and S. robusta Horik. (= S. subnimbosa Steph.) (Amakawa & Hattori, 1953; Inoue, 1972), but seem to have been confused with paraphyses because the male bracts are usually extremely similar to vegetative leaves and they are very difficult to recognise, especially when the antheridia shrivel in herbar-The terms paraphyllia ium specimens. and pseudoparaphyllia have also been used interchangeably in earlier reports (*e.g.* Cao *et al.*, 2004; Cao & Sun, 2008). It is still necessary to investigate the occurrence of the true paraphyllia, and evaluate their taxonomic value in *Scapania*.

Scapania koponenii may be confused with the European S. aspera. The former, however, differs mainly in its smaller size (1.2-2 mm wide and 5–20 mm long), smaller leaf cells (median $13-16 \times$ $16-20 \,\mu\text{m}$), dense \pm hemispherical papillae, and the more elongated, \pm spinose terminal tooth cells in the leaf margins (Potemkin, 2000; Borovichev et al., 2016). The molecular phylogenetic study also reveals the two species belong to different main clades of Scapania (Heinrichs et al., 2012). In China, S. koponenii is most similar to S. ciliata, but in the latter the plants are 2-4 cm long, with marginal teeth of the leaf often unicellular, rather strongly elongated and notably spinose, dense, and hyaline, leaf lobes usually 1/2 the leaf lobule in size, and perianth mouth with cilia 1-4 cells long.

Taxonomic treatment

Scapania koponenii Potemkin, Ann. Bot. Fenn. 37 (1): 41. 2000. (Figures 2–3)

Type: China, Hunan Prov.: Yizhang Co., Mt. Mangshan, Guizizhai, core area of the forest reserve. Primeval subtropical (warm temperate) *Cyclobalanopsis* Oerst., *Lithocarpus* Blume, *Pinus kwangtungensis* Chun & Tsiang, *Rhododendron* L, *Schisma* Dumort. forest on slope, 24°57′N, 112°55′E, 1160 m, on cliff on open moist slope, 2 Oct. 1997, *T. Koponen, S. Huttunen* & *P.-C. Rao* 50767*a* (holotype: H; isotypes: LE and herb. Forest Botanical Garden, Changsha).

= Scapania paraphyllia T.Cao, C.Gao, J.Sun & B.R.Zuo, Acta Phytotax. Sin. 45 (3): 311. 2007. syn. nov. Type: China, Zhejiang Prov.: Suichang Co., Mt. Jiulong, 1360 m, on rock, 23 Apr. 1981, Z. L. Liu 554 (holotype: HSNU!).

Representative specimens examined: China: Fujian, Wuyishan Co., Wuyishan National Nature Reserve, 27°44′59.57″N, 117°40′46.10″E, 695 m, on rock with a thin layer of soil, 31 Aug. 2015, Zhang et al. 20150831-101 (HSNU), ibid., 27°44′59.96″N, 117° 40'46.37"E, 695 m, on rock with a thin layer of soil, 31 Aug. 2015, Zhang et al. 20150831-107 (HSNU), ibid., 27°43′58.85″N, 117°38′44.19″E, 1061 m, on rock with a thin layer of soil, 30 Aug. 2015, Zhang et al. 20150830-145 (HSNU), ibid., 27°43'58.96"N, 117°39'22.83"E, 937 m, on rock, 30 Aug. 2015, Zhang et al. 20150830-178, 20150830-178A (HSNU); Guangdong, Ruyuan Co., Babaoshan Nature Reserve, 800-900 m, on rock, 2 Nov. 1995, Li et al. 95233 (paratype of S. koponenii: HSNU!, SYS!); Xingan Longtangjiang, 25° Guangxi, Co., 48'43.61"N, 110°25'48.55"E, 446 m, 10 Jul. 2015,

Wei et al. 20150710-3 (HSNU); Guizhou, Leishan Co., Leigongshan National Nature Reserve, 26° 22.163'N, 108°08.994'E, 964 m, on rock, 26 Aug. 2010, Zhu et al. 20100826-92B (HSNU); Jiangxi, Jinggangshan Co., Jinggangshan National Nature Reserve, on rock with a thin layer of soil, 15 Oct. Wang 20141015-13 (HSNU), Zixi Co., 2014. Matoushan National Nature Reserve, 27°47′58.63″N, 117°12'07.93"E, 438 m, on wet cliff, 27 Jul. 2015, Zhang 20150727-52 (HSNU), Yanshan Co.. Wuyishan National Nature Reserve, 27°50'17.39"N, 117°44'32.82"E, 1532 m, on wet cliff, 3 Aug. 2015, Zhang 20150803-137 (HSNU), ibid., 27°50'26.46"N, 117°43′55.69″E, 905 m, on rock with a thin layer of soil, 4 Aug. 2015, Zhang 20150804-76 (HSNU); Zhejiang, Longquan City, Fengyangshan National Nature Reserve, 27°54.444'N, 119°10.452'E, 1293 m, on rock, 19 Apr. 2011, Zhu & Wei 20110419-12 (HSNU), Suichang Co., Mt. Jiulong, on rock, 1360 m, 23 Apr. 1981, Z. L. Liu 553 (paratype of Scapania paraphyllia: HSNU!). Taishun Co., Wuyanling National Nature Reserve, on wet rock, 28 Jul. 2001, Cao 010281 (SHTU).

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Taxonomic Additions and Changes: *Scapania koponenii* Potemkin (*S. paraphyllia* T.Cao, C.Gao, J.Sun & B.R.Zuo *syn. nov.*).

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Appendix 1

Voucher information and GenBank accession numbers for the taxa used for this study. Sequences newly generated are in bold. A long dash indicates missing sequences.

		GenBank accession number		
Taxon	Locality and voucher	nrITS	<i>trn</i> L-F	atpB- rbcL
Scapania aequiloba (Schwagr.) Dumort	Germany, Thuringia, Grosburschla, Marstaller 2-7-2001 (JE)	JN631360	JN631498	JN631629
Scapania curta (Mart.)	Germany, Saxony-Anhalt, Breitenbach, Hentschel Bryo 3174	JN631404	JN631542	JN631671
Scapania americana Mull.Frib.	U.S.A., Oregon, Douglas Co., <i>Shevock 26373</i> (GOET)	JN631367	JN631505	JN631636
Scapania americana	U.S.A., California, San Mateo Co., Shevock 27851 (GOET)	JN631365	JN631503	JN631634
Scapania americana	U.S.A., California, San Mateo Co., Shevock 27855 (GOET)	JN631366	JN631504	JN631635
<i>Scapania ciliata</i> Sande Lac.	Nepal, Kangchenjunga, Sikdim — Chauki, <i>Long 17560</i> (JE)	JN631393	JN631530	JN631661
Scapania ciliata	China, Guizhou, Yuao, <i>Peng 20100518-13</i> (HSNU)	JN631391	JN631528	JN631659
Scapania ciliata	China, Guangxi, Laibin, Ye & Wei 20090719-10 (HSNU)	JN631390	JN631527	JN631658
(Roth) Dumort.	Germany, Saxony Anhalt, Ireseburg-Ihale, <i>Eckstein 1409</i> (GOET)	JN631398	JN631534	JN631663
Scapania compacta	Spain, La Palma, Cubo de Galga, Huneck JE-H3294 (JE)	JN631399	JN631537	JN631666
Scapania compacta	United Kingdom, Argyll, Glencoe, Long & Murray 11492 (JE)	JIN631400	JN631538	JN631667
Scapania griffitnii Schiffn.	China, Fujian, Denua Co., Zhu et al. 20100403-21 (HSNU)	JIN631419	JIN631557	JIN631684
Scapania griniumi Scapania iavanica	Indonesia, Control Sulawesi, Mt. Pereketimbu, Gradetain 12046	JIN031420	JIN03 1000	JIN03 1003
Gottsche	(GOET)	JIN03 1433	JIN03 137 1	JIN03 1097
Scapania javanica	Indonesia, Sumatra, Berastagi, <i>Schafer-Verwimp & Verwimp</i> 24861 (GOET)	JN631436	JN631572	JN631698
<i>Scapania kaurinii</i> Ryan	Russia, Chita Prov., <i>Bakalin 11-1-00</i> (KPABG)	EU791759	EU791650	
<i>Scapania koponenii</i> Potemkin	China, Zhejiang, Suichang Co., Zhu et al. 20090630-22 (HSNU)	JN631438	JN631574	JN631700
Scapania koponenii	China, Zhejiang, Suichang Co., Zhu et al. 20090630-48 (HSNU)	JN631437	JN631573	JN631699
Scapania koponenii	China, Fujian I, Wuyishan Co., <i>Zhang et al. 20150830-178A</i> (HSNU)	KY311839	KY311849	KY311860
Scapania koponenii	China, Fujian II, Wuyishan Co., <i>Zhang et al. 20150830-145</i> (HSNU)	KY311840	KY311850	KY311861
Scapania koponenii	China, Fujian III, Wuyishan Co., <i>Zhang et al. 20150831-107</i> (HSNU)	KY311841	KY311851	KY311862
Scapania koponenii	China, Jiangxi I, Yanshan Co., <i>Zhang 20150804-76</i> (HSNU)	KY311842	KY311852	KY311863
Scapania koponenii	China, Jiangxi II, Jinggangshan Co., <i>Wang 20141015-13</i> (HSNU)	KY311843	KY311853	KY311864
Scapania koponenii	China, Jiangxi III, Zixi Co., Zhang 20150727-52 (HSNU)	KY311844	KY311854	KY311865
Scapania koponenii	China, Guangxi, Xingan Co., Wei et al. 20150/10-3 (HSNU)		KY311855	KY311866
Scapania ligulata Steph.	North Korea, Kumgangsan, Manmulsang, <i>Huneck KDVR 88-36</i>	JN631439 JN631443	JN631575 JN631579	 JN631704
Scapania ligulata	Nepal Kanacheniunga Nesum-Buie Daurali <i>Long</i> 17499 (JE)	.IN631442	JN631578	JN631703
Scapania ligulata	China, Guangxi, Maozhou, <i>Wei 20090705124</i> (HSNU)	JN631440	JN631576	JN631701
Scapania ligulata	China, Fujian, Dehua Co., <i>Zhu et al. 20100406-1B</i> (HSNU)	JN631441	JN631577	JN631702
Scapania ligulata	China, Jiangxi, Yanshan Co., Zhang 20150803-120A (HSNU)	KY311845	KY311856	KY311867
Scapania paraphyllia T.Cao, C.Gao, J.Sun &	China, Jiangxi, Yanshan Co., <i>Zhang 20150803-137</i> (HSNU)	KY311846	KY311857	KY311868
Scapania paraphyllia	China, Fujian, Wuyishan Co., <i>Zhang et al. 20150830-178</i> (HSNLI)	KY311847	KY311858	KY311869
Scapania sphaerifera H Buch & Tuom	(KPARG) (KPARG)	EU791765	EU791656	
Scapania sphaerifera	Russia, Siberia, Buryatiya Rep., <i>Konstantinova, Hep. Ross. Exs.</i> 20 (JE)	JN631471	JN631605	JN631730
<i>Scapania spitsbergensis</i> (Lindb.) Mull.Frib.	Norway, Spitsbergen, Konstantinova 90-2-06 (KPABG)	EU791761	EU791752	—
Scapania verrucosa Heeg	Russia, Caucasus, Karachayevo-Cherkessian Rep., Konstantinova 609/6-05 (KPABG)	EU791763	EU791654	—
Scapania verrucosa	China, Xizang, Linzhi Pre., Wang et al. 20140826-21 (HSNU)	KY311848	KY311859	KY311870